

PTO/PCT Rec'd 10 JAN 2002

Method For Preparing Cyclohexane Carboxylic AcidsArea of the Invention

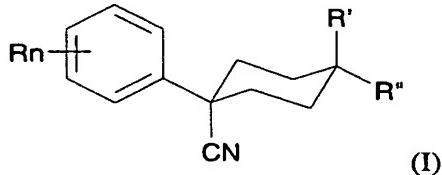
This invention relates to a method for preparing 4-substituted-4-cyanocyclohexanecarboxylic acids. Exemplary compounds are useful as PDE 4 inhibitors.

5 Background of the Invention

The process of this invention relates to making compounds which are useful in treating diseases modulated by the isoforms of the phosphodiesterase 4 enzyme. The novel intermediates and processes of this invention are useful in making acids which are known PDE 4 inhibitors. They are useful for, among other things, treating pulmonary diseases such 10 as chronic obstructive pulmonary disease (COPD) and asthma. The compounds which are prepared by the methods of this invention are described in, for example U.S. patent 5,554,238 issued 03 September, 1996. That patent is incorporated here by reference in full. Those compounds, particularly the 4-cyanocyclohexanoic acids, have marked effects on neutrophil activity, inhibiting neutrophil chemotaxis and degranulation *in vitro*. In animal 15 models, those compounds reduce neutrophil extravasation from the circulation, pulmonary sequestration and the edematous responses to a number inflammatory insults *in vivo*. They have been found to be useful in treating COPD in humans, and possibly in other mammalian species which suffer from COPD.

Summary of the Invention

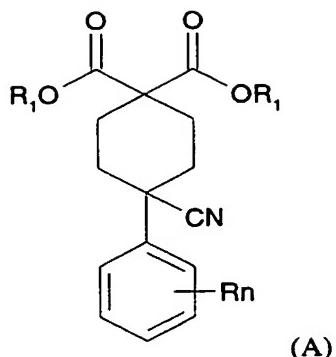
20 In a first aspect, this invention relates to a process for preparing a compound of formula (I)



where

25 R is halo, C₁-6alkyl, C₁-6alkyl substituted with 1 to 4 halogens, C₁-6alkoxy, C₁-6alkenyl, -O-(CH₂)_mcycloalkyl of 3-6 carbons;
 n is 1-5;
 m is 0 - 6; and
 R' and R'' are independently hydrogen or CO(O)X where X is hydrogen or C₁-30 6alkyl;

which process comprises decarboxylating the diester or diacid of Formula (A)



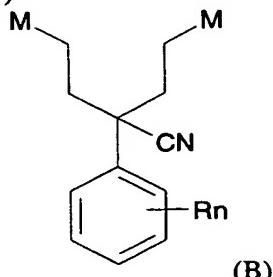
(A)

wherein R_1 is hydrogen or a C_{1-6} -alkyl-ester forming group and R and n are the same as for Formula (I).

In a further aspect this invention relates to a compound of formula (A) per se.

- 5 In a third aspect this invention relates to preparing certain other intermediates that are useful in preparing the diester or di-acid of Formula (A), and the intermediates themselves, i.e:

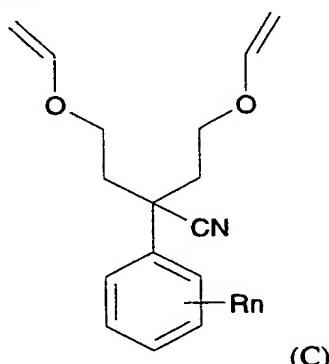
a compound of Formula (B)



(B)

- 10 wherein R and n are the same as in Formula (I) and M is OH , an activated hydroxyl group, or halo; and

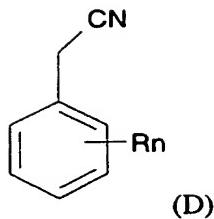
a compound of Formula (C)



(C)

wherein R and n are the same as in Formula (I).

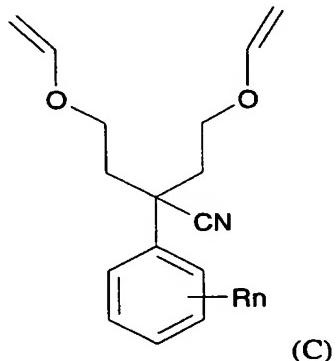
- 15 In yet another aspect, the invention provides a method for making a compound of Formula (C) by treating the nitrile of formula (D)



where R and n are the same as defined above, with 2-chloroethyl vinyl ether and a strong base.

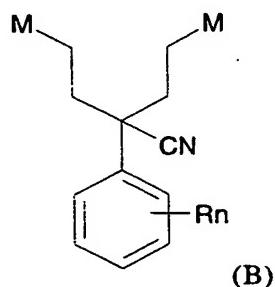
This invention also provides a method for preparing a compound of Formula (I)
5 which comprises

- a. converting the vinylethyl ether of Formula (C)



10 wherein R and n are halo, C₁₋₆alkyl, C₁₋₆alkyl substituted with 1 to 4 halogens, C₁₋₆alkoxy, C₁₋₆alkenyl, -O-(CH₂)_mcycloalkyl of 3-6 carbons;
n is 1-5;
m is 0 - 6;
to a compound of Formula (B)

15

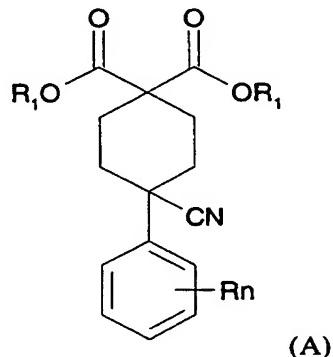


where M is OH,

b. converting the hydroxyl group of Formula (B) to a compound of Formula (B)
20 where M is a tosylate, mesylate or a triflate,

- c. converting the tosylate, mesylate or triflate in Formula (B) to a compound of Formula (B) where M is halo,
- d. treating the di-halo compound with dialkyl malonate to obtain a compound of Formula (A)

5



where R_1 is lower alkyl,

- e. optionally saponifying the diester of Formula (A) to obtain a compound of Formula (A) where R_1 is hydrogen; and
- 10 f. decarboxylating a compound of Formula (A) where R_1 is hydrogen or C₁-6alkyl to obtain a compound for Formula (I) where one of R' is hydrogen and the other is CO(O)X where X is C₁-6alkyl or hydrogen.

Specific Embodiments of the Invention

This invention provides a method for preparing cyclohexanoic acids. In particular it provides an alternative means for preparing the cyclohexanoic acids disclosed in U.S. patent 5,554,238 where the 4-position on the cyclohexane ring has a CN group.

"Halo" as used herein includes fluoro, chloro, bromo, and iodo. "Halide" includes fluoride, chloride, bromide and iodide.

For all of the compounds disclosed herein, a preferred embodiment is one where there are two R groups, i.e. n is 2. Most preferred are those compounds where one R group is at the 3 position and the second R group is on the 4 position of the benzene ring. More particularly it is preferred that each R group be independently C₄-6cycloalkyloxy or C₁-2alkoxy unsubstituted or substituted by 1 or more halogens. More preferred are methoxy, C₁-2alkoxy substituted by up to 3 fluoro atoms, cyclopropylmethoxy or cyclopentyloxy.

25 The more preferred R groups are those wherein the 4-position R group is methoxy, -O-CF₃, -O-CHF₂, or -O-CH₂CHF₂. and the 3-position R group is cyclopropylmethoxy or cyclopentyloxy.

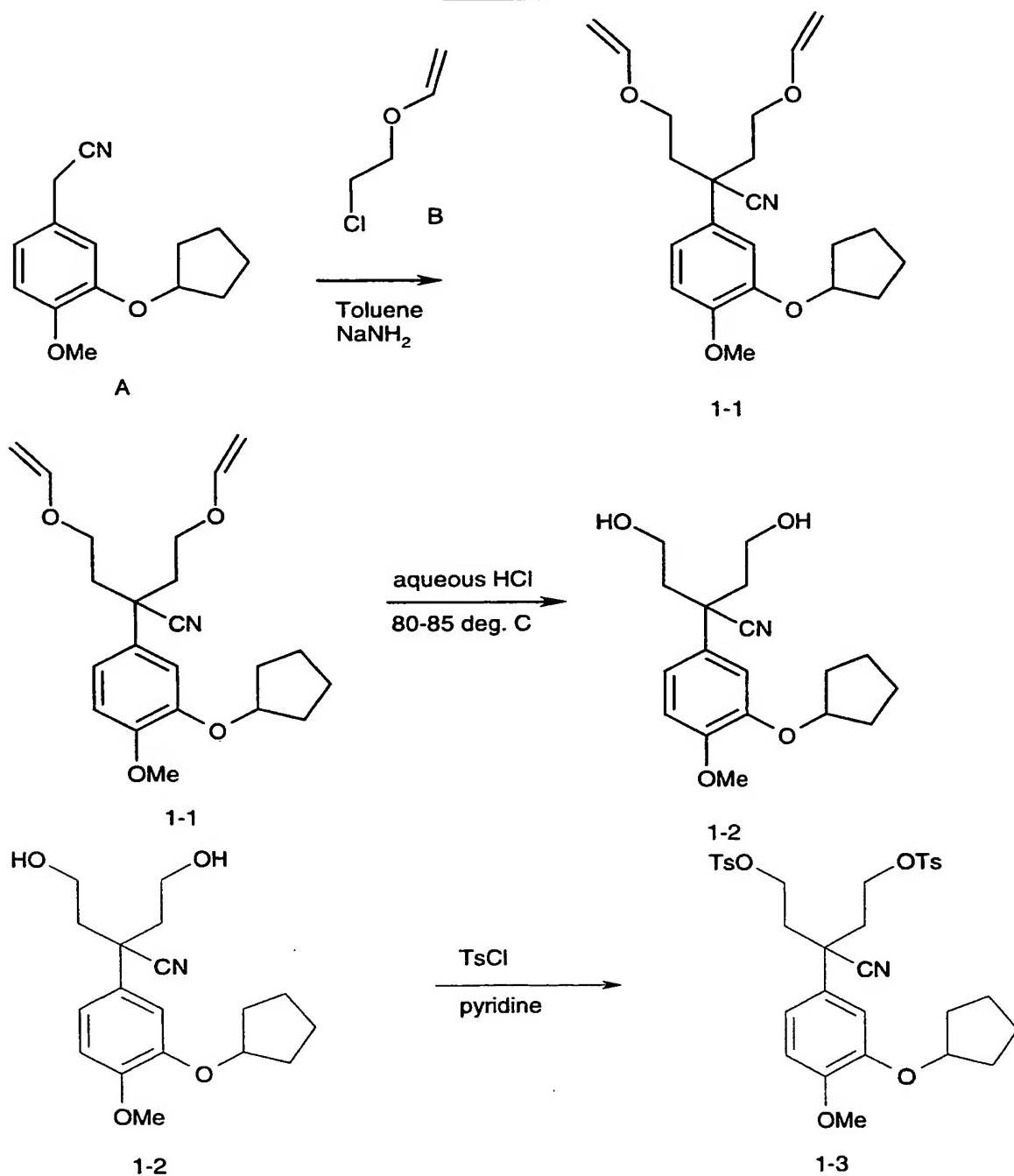
In Formula (A) the most preferred R_1 groups are hydrogen, methyl or ethyl.

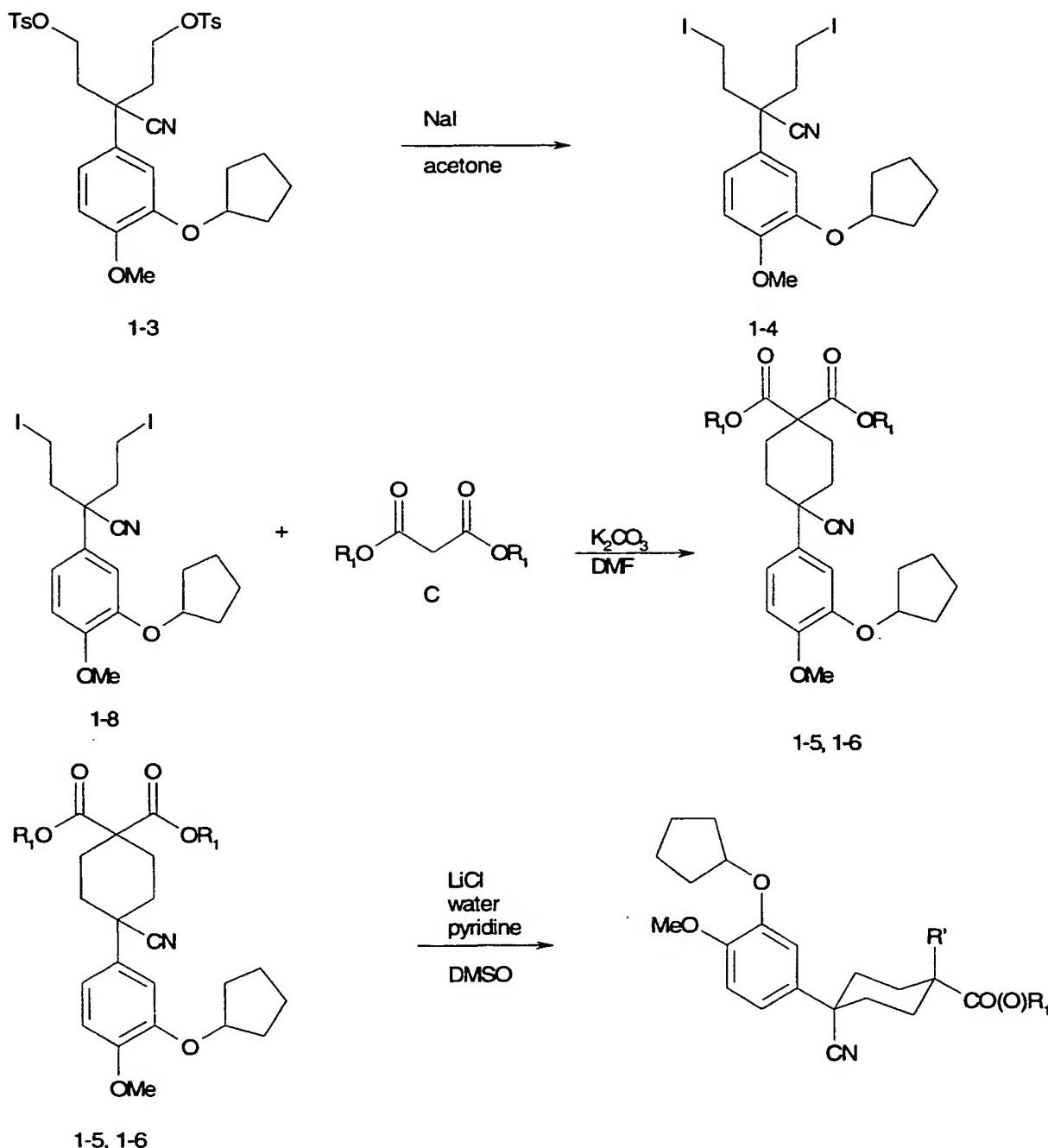
In Formula (B) the most preferred M groups are OH, tosyl and iodo.

The most preferred product of the process of this invention are those compounds which have a 3-cyclopentyloxy-4-methoxyphenyl substitution pattern.

Reaction Scheme I provides a diagrammatic overview of the intermediates and chemistries employed in this invention.

5

Scheme I



- 5 The starting material 3-cyclopentyloxy-4-methoxybenzenecetonitrile is a known compound. See for example U.S. patent 5,449,686. The 2-chloroethylvinyl ether is commercially available (Aldrich). To effect the reaction, a strong base is charged to a reaction vessel containing a suitable non-polar solvent to which the vinyl ether is added. This mixture is heated to between about 30 to 70 °C and charged with the
- 10 benzeneacetonitrile (A) pre-dissolved in the same solvent as the base and the vinyl ether.

Toluene is a preferred solvent. A preferred base is sodium amide. The amount of base is equivalent, on a molar basis, to that of the vinyl ether. Both are used in about a three-fold excess relative to the substrate. After the benzeneacetonitrile has been charged to the reaction flask, the solution is further heated to around 80 °C more or less. Usually the 5 reaction is complete in about 30 minutes to 2 hours. The product (1-1) is isolated using standard procedures.

The bis 2-hydroxyethyl compound (1-2) is prepared by treating the vinyl ether moiety prepared as per the preceding paragraph with a strong mineral acid in an aqueous solvent. For example water can be added to the 2-(ethenyloxy)ethyl compound (1-1), 10 heating that combination to about 70-90 °C and then adding a molar excess of a mineral acid such as HCl or the like. A preferred set of conditions is one where the 2-(ethenyloxy)ethyl is treated with water and heated to about 80 °C more or less followed by the addition of a 50% molar excess of concentrated HCl. Under these conditions the reaction is complete in 5 - 20 minutes.

15 To obtain the halogenated compound 1-4, the diol is converted to a group which can be displaced by a halide ion. For example the diol can be converted to a tosylate, mesylate, or the like, by treating the diol with reagents and under conditions which form the tosylate, etc. By way of example the diol is dissolved in an organic solvent and treated with an excess of *p*-toluenesulfonyl chloride at room temperature for 3- 7 hours. Preferably the 20 reaction is run in pyridine with about a 2.5 molar excess of the *p*-toluenesulfonyl chloride.

This tosylate (or mesylate, triflate, etc) (1-3) is converted to the di-halo 1-4 by dissolving it in a polar aprotic solvent, and adding a weak base and a halide salt. This mixture is heated to reflux for a number of hours, for example overnight. A preferred solvent is acetone or dimethyl formamide. A preferred halide salt is sodium or lithium 25 iodide though other sodium or potassium salts of fluorine, chlorine and bromine can be used as well. A 2 to 6-fold excess of the halide salt is preferred. Refluxing overnight usually effects completion of the reaction.

Forming the cyclohexane dicarboxylates or diacids 1-5 and 1-6 is effected by charging the di-halo compound (1-4) to a solution of a dialkyl malonate or malonic acid and 30 a weak base in a dipolar aprotic solvent. This slurry is stirred for an extended period of time at an elevated temperature, for example overnight. More specifically sodium or potassium carbonate is combined with the likes of dimethyl malonate in a solvent such as dimethylformamide. Then the di-halo 1-4 is added and the resulting slurry is stirred 35 overnight at about 75-95 °C or so. The malonate is added in about a 1:1 molar ratio to that of the di-halo compound.

The diester may be saponified to give the diacid, though this step is not illustrated in Scheme 1. This is accomplished by treating the diester with an aqueous base in a water-

miscible solvent. For example the diester is charged to a reaction vessel containing the likes of tetrahydrofuran to which is added water and an alkali hydroxide base such as lithium hydroxide. This solution is heated at reflux for a number of hours, for example overnight.

Decarboxylating the diester or dicaid to get the mono-ester or mono-acid is
5 accomplished by dissolving the diester in the likes of dimethylsulfoxide, adding about an equivalent of a base such as pyridine, about 3 equivalents of water and about 3 equivalents of a salt such as lithium chloride. This solution is stirred for several hours at 100 to 150 °C or thereabouts for 4-8 hours. Product is extracted from an acidified aqueous solution and further purified by conventional means. The product is a mixture of cis and trans isomers in
10 about a 1:1 ratio. The *cis* form of the ester or acid can be enriched by dissolving a mixture of isomers in a lower alkanol and treating that solution with the alkali metal salt of the alkanol. A preferred alkanol is *t*-butanol and a preferred alkali metal salt is potassium *t*-butanol. The acid may be obtained by saponifying the ester using a base and then acidifying the resulting salt with using a mineral acid, for example.

15 The following examples are provided to illustrate the invention. These illustrative examples are not intended to limit the claimed invention in any fashion.

Examples

Example 1

Preparation of 3-(Cyclopentyloxy)- α,α -bis[2-(ethenyloxy)ethyl]-4-methoxybenzeneacetonitrile

20 A 1 L flask was charged with 150 mL of toluene, sodium amide (16.5 g, 0.38 mole, 2.9 equivalents), and 2-chloroethyl vinyl ether (41.9 g, 0.39 mole, 3.0 equivalents). The suspension was heated to 50 °C, then charged with a solution of 3-cyclopentyloxy-4-methoxybenzeneacetonitrile (30 g, 0.13 mole, 1.0 equivalents) in 150 mL of toluene. The 25 reaction was then carefully heated to 80 °C. The progress was followed by HPLC (acetonitrile/0.1 N aqueous ammonium acetate at 65/35, 15 cm Beckman ODS Ultrasphere, 2 mL/min, 215 nm UV). After 60 minutes, the solution was poured into 1 L of water and 300 mL of *t*-butyl methyl ether. The layers were separated, the organic layer washed with water, then brine. The solvent was removed under reduced pressure to give a brown oil
30 (52.1 g). The captioned compound was isolated by column chromatography (230-400 mesh silica gel, 10/1 hexane/ethyl acetate).

Mass spectrometry gave m/z = 372 (M+H⁺)⁺

(¹H NMR, 300 MHz, CDCl₃, δ ppm) δ 1.55-1.65 (m, 2H, ring CH₂), δ 1.70-2.00 (m, 6H, ring CH₂'s), δ 2.2-3.5 (m, 4 H,(CH₂)₂CCN), δ 3.50-3.85 (m, 4H, CH₂O), δ 3.85 (s, 3H, OCH₃), δ 3.95-4.10 (m, 4H, CH₂ alkene), δ 4.8 (m, 1H, ring CH), δ 6.30-6.40 (m, CH, alkene), δ 6.85-7.0 (m, 3H, aromatic)

Example 2Preparation of 3-(Cyclopentyloxy)- α,α -bis(2-hydroxyethyl)-4-methoxybenzenecetonitrile

Purified 3-(cyclopentyloxy)- α,α -bis[2-(ethenyloxy)ethyl]-4-methoxybenzenecetonitrile (5 g, 13.5 mmol) was treated with water (50 mL) and heated to 80 °C with rapid stirring. Concentrated hydrochloric acid (1.85 mL, 22.2 mmol) was added and stirring was continued for 10 minutes. The solution was poured into ice water (50 mL) and methylene chloride (50 mL). The layers were separated, and the aqueous layer was extracted once with methylene chloride. The combined organic layers were washed with water, then brine, and concentrated to a light yellow oil in quantitative yield. Structure and purity were confirmed by ^1H NMR.

(^1H NMR, 300 MHz, CDCl_3 , δ ppm) δ 1.55-1.70 (m, 2H, ring CH_2), δ 1.75-1.95 (m, 6H, ring CH_2 's), δ 2.10-2.40 (m, 4H, $(\text{CH}_2)_2\text{CCN}$), δ 3.55-3.85 (m, 4H, CH_2O), 3.88 (s, 3H, OMe), δ 4.8 (m, 1H, ring CH), δ 6.80-7.00 (m, 3H, aromatic)

Example 3

15 Preparation of 3-(Cyclopentyloxy)-4-methoxy- α,α -bis[2-[(4-methylphenyl)sulfonyl]oxy]ethyl]benzenecetonitrile

A 250 mL flask was charged with 3-(cyclopentyloxy)- α,α -bis(2-hydroxyethyl)-4-methoxybenzenecetonitrile (6.4 g, 20.1 mmol), pyridine (65 mL), and *p*-toluenesulfonyl chloride (9.56 g, 50.2 mmol). The solution warmed slightly (exothermic), then was stirred at room temperature for 5 hours. The reaction was deemed complete by HPLC (acetonitrile/0.1 N aqueous ammonium acetate at 65/35, 15 cm Beckman ODS Ultrasphere, 2 mL/min, 215 nm UV). The reaction was poured into 100 mL of 5% HCl and 50 mL of methylene chloride. The layers were separated, and the organic layer was washed with 5% HCl until neutral. The neutralized organic layer was then washed once with brine and concentrated. The captioned compound was isolated as a white solid by crystallization from ethanol and t-butyl methyl ether.

Mass spectrometry gave $m/z = 645 (\text{M}+\text{NH}_4^+)^+$

(^1H NMR, 300 MHz, CDCl_3 , δ ppm) δ 1.55-1.70 (m, 2H, ring CH_2), δ 1.72-2.0 (m, 6H, ring CH_2 's), δ 2.20-2.45 (m, 4H, $(\text{CH}_2)_2\text{CCN}$), 2.45 (s, 3H, ar- CH_3), 3.85 (s, 3H, OMe), δ 3.85-4.28 (m, 4H, CH_2O), δ 4.75 (m, 1H, ring CH), δ 6.75-7.75 (11H indicated, aromatic)

Example 4Preparation of 3-(Cyclopentyloxy)- α,α -bis(2-iodoethyl)-4-methoxybenzenecetonitrile

A 250 mL flask was charged with 3-(cyclopentyloxy)-4-methoxy- α,α -bis[2-[(4-methylphenyl)sulfonyl]oxy]ethyl]benzenecetonitrile (5.0 g, 7.97 mmol), acetone (75 mL), and sodium bicarbonate (50 mg). This solution was stirred well while sodium iodide (5.98 g, 39.9 mmol) was added, then heated to reflux overnight. The reaction was poured into

aqueous ammonium chloride and sodium bisulfite, then extracted with *t*-butyl methyl ether. The organic layer was dried over sodium sulfate, then concentrated to a clear colorless oil. The captioned compound was crystallized from *t*-butyl methyl ether and hexanes to give a white solid.:.

5 Mass spectrometry gave m/z = 540 (M+H⁺)⁺

CHN analysis calculated for C₁₈H₂₃NO₂I₂ (539.12): C 40.10, H 4.30, N 2.60; found: C 40.06, H 4.30, N, 2.45.

(¹H NMR, 300 MHz, CDCl₃, δ ppm) δ 1.55-1.70 (m, 2H, ring CH₂), δ 1.75-2.05 (m, 6H, ring CH₂'s), δ 2.35-2.62 (m, 4H, (CH₂)₂CCN), δ 2.8 (m, 2H, CH₂I), δ 3.15 (m,

10 2H, CH₂I), 3.87 (s, 3H, OMe), δ 4.8 (m, 1H, ring CH), δ 6.85 (s, 1H, aromatic), δ 7.18-7.30 (m, 2H, aromatic)

Example 5

Preparation of Dimethyl 4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]-1,1-cyclohexanedicarboxylate and Diethyl 4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]-1,1-cyclohexanedicarboxylate

15 A 125 mL flask was charged with 60 mL of dimethylformamide, potassium carbonate (5.02 g, 36.4 mmol, 3.6 equivalents), dimethylmalonate (1.33 g, 10.1 mmol, 1.0 equivalents), and then 3-(cyclopentyloxy)-α,α-bis(2-iodoethyl)-4-methoxybenzeneacetonitrile (6.0 g, 11.1 mmol, 1.1 equivalents). The slurry was stirred at 80 °C overnight. The completed reaction 20 was poured into 50 mL of water and 50 mL of *t*-butyl methyl ether. The organic layer was extracted three times with water, then once with brine. The product was isolated by column chromatography (Flash silica [230-400 mesh], 80/20 hexanes/ethyl acetate). Alternatively it was crystallized from hexanes/ethyl acetate (3/1) to give white crystals. The diethyl derivative was prepared using the same procedure.

25 Dimethyl ester:

Mass spectrometry gave m/z = 416 (M+H⁺)⁺

CHN analysis calculated for C₂₃H₂₉NO₆ (539.12): C 66.49, H 7.04, N 3.37; found: C 66.24, H 6.94, N, 3.33.

(¹H NMR, 300 MHz, CDCl₃, δ ppm) δ 1.55-1.67 (m, 2H, ring CH₂), δ 1.75-2.60 (m, 14H, ring CH₂'s), 3.75 (s, 3H, CO₂Me), 3.78 (s, 3H, CO₂Me), 3.82 (s, 3H, OMe), δ 4.8 (m, 1H, ring CH), δ 6.80-7.02 (m, 3H, aromatic)

Diethyl ester

Mass spectrometry gave m/z = 444 (M+H⁺)⁺

mp. 74.0-74.5

35 (¹H NMR, 300 MHz, CDCl₃, δ ppm) δ 1.20-1.38 (m, 6H, ethyl CH₃), δ 1.50-2.60 (m, 16H, ring CH₂'s), 3.85 (s, 3H, OMe), δ 4.65-4.85 (m, 4H, ethyl CH₂), δ 4.8 (m, 1H, ring CH), δ 6.80-7.05 (m, 3H, aromatic).

Example 6Hydrolysis of the diethyl ester to the diacid

A 25 mL flask was charged with tetrahydrofuran (5 mL), the diethyl ester (SB 220523, 0.5 g, 1.13 mmol, 1.0 equivalent), water (5 mL), and lithium hydroxide monohydrate (0.95 g, 22.6 mmol, 20 equivalents). The solution was stirred at reflux for 18 hours. The reaction was deemed complete by HPLC (15 cm Supelcocol LC-ABZ, 40/60/0.1 [acetonitrile/water/TFA], 1.5 mL/min., 215 nm UV). The reaction solution was then cooled and diluted with 10% HCl and t-butyl methyl ether. The layers were separated and the aqueous layer was washed once with t-butyl methyl ether. The organic layers were combined and washed with water and then brine. The solution was then concentrated to a tan solid. Water was removed by reconcentrating once with acetonitrile. The crude product was obtained in about 90% yield. The crude product showed residual ethyl ester (< 5%) by HPLC and ^1H NMR.

(^1H NMR, 300 MHz, CDCl_3 , δ ppm) δ 1.55-1.70 (m, 2H, ring CH_2), δ 1.75-2.35 (m, 12H, ring CH_2 's), δ 2.52-2.63 (m, 2H, ring CH_2), 3.85 (s, 3H, OMe), δ 4.8 (m, 1H, ring CH), δ 6.8-7.0 (m, 3H, aromatic)

Example 7Decarboxylation of the diester: Ethyl 4-cyano-4-(3-cyclopentyloxy-4-methoxyphenyl)-cyclohexanecarboxylate

A 100 mL flask was charged with dimethylsulfoxide (35 mL), dimethyl 4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]-1,1-cyclohexanedicarboxylate (3.5 g, 8.43 mmol, 1.0 equivalent), water (0.455 g, 25.3 mmol, 3.0 equivalents), pyridine (0.66 g, 8.43 mmol, 1.0 equivalent), and lithium chloride (1.07 g, 25.3 mmol, 3.0 equivalents). The solution was stirred at 130 °C for 6.5 hours. The reaction solution was then cooled and diluted with 1% HCl and t-butyl methyl ether. The layers were separated and the organic layer was washed with water twice and with brine once. The solution was concentrated to a clear oil. Water was removed by reconcentrating once with methanol. The product was obtained in quantitative yield as a clear oil and as a mixture of *cis* and *trans* isomers in about a 1:1 ratio.

Mass spectrometry gave m/z = 372 ($\text{M}+\text{H}^+$)⁺

(^1H NMR, 300 MHz, CDCl_3 , δ ppm) δ 1.29 (t, 3H, ethyl CH_3), δ 1.55-1.70 (m, 2H, ring CH_2), δ 1.75-2.30 (m, 14H, ring CH_2 's), δ 2.75-2.80 (m, 1H, CHCO_2Et), 3.85 (s, 3H, OMe), δ 4.13-4.22 (q, 2H, ethyl CH_2), δ 4.8 (m, 1H, ring CH), δ 6.8-7.0 (m, 3H, aromatic)

Example 8Preparation of 4-cyano-4-(3-cyclopentyloxy-4-methoxyphenyl)-cyclohexanecarboxylic acid

The isomeric mixture (approximately 1 to 1 ratio) of the methyl esters prepared in Example 5 (2.94 g, 8.2 mmol, 1.0 equivalent) was dissolved in *t*-butanol (30 mL) under a nitrogen atmosphere. Potassium *t*-butoxide (1.8 g, 16.5 mmol, 2.0 equivalent) was added

and the mixture was stirred 6-18 hours to give a ratio of *cis* to *trans* isomers of 14 to 1. The same procedure was used to treat the ethyl esters and gave a ratio of 8 to 1. The ratios were monitored using HPLC (15 cm Supelcocol LC-ABZ, 40/60/0.1 [acetonitrile/water/TFA], 1.5 mL/min., 215 nm UV).

- 5 To hydrolyze the equilibrated ester product, two drops of water were added to the reaction solution and the solution was stirred until no ester could be detected. The reaction was then diluted with *t*-butyl methyl ether and 5% HCl (the pH of the aqueous layer was between 1-2). The layers were separated and the organic layer was washed with brine. The ratio of *cis* to *trans* acid was improved even further (to 121 to 1) by crystallizing the
10 *cis/trans* mixture from 20 ml of hexanes/ethyl acetate (3/1)

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